L Number	Hits		DB	Time stamp
1	9106	serine adj (protease or proteinase)	USPAT;	2003/11/18 10:53
			US-PGPUB;	
			EPO; JPO;	
_			DERWENT	
2	9630		USPAT;	2003/11/18 10:56
		Fluorescent or fret)	US-PGPUB;	
			EPO; JPO;	
	600		DERWENT	0000 /11 /10 10 56
3	633	,, ·, ·1, ·, ·	USPAT;	2003/11/18 10:56
		(substrate near4 (Fluorescence or	US-PGPUB;	
		Fluorescent or fret))	EPO; JPO; DERWENT	
4	0	((serine adj (protease or proteinase)) and	USPAT;	2003/11/18 10:58
4	U	(substrate near4 (Fluorescence or	US-PGPUB;	2003/11/18 10:38
		Fluorescent or fret)	EPO: JPO:	
		adj energy and transfer)	DERWENT	
5	63	((serine adj (protease or proteinase)) and	USPAT:	2003/11/18 11:01
	03	(substrate near4 (Fluorescence or	US-PGPUB;	2003/11/10 11:01
		Fluorescent or fret))) and (fluorescence	EPO; JPO;	
		adj resonance adj energy adj.transfer)	DERWENT	
6	54		USPAT;	2003/11/18 11:02
_	· ·	((fluorescence adj resonance adj energy	US-PGPUB;	= = = = = = = = = =
		adj transfer) or fret)	EPO; JPO;	
ł			DERWENT	
7	6	((proteinase or protease) near5	USPAT;	2003/11/18 11:02
		((fluorescence adj resonance adj energy	US-PGPUB;	
		adj transfer) or fret)) and ((serine adj	EPO; JPO;	
		(protease or proteinase)) and (substrate	DERWENT	
		near4 (Fluorescence or Fluorescent or		
i		fret)))		

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                 present
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     4 AUG 05
                 New pricing for EUROPATFULL and PCTFULL effective
                 August 1, 2003
NEWS
     5 AUG 13
                 Field Availability (/FA) field enhanced in BEILSTEIN
     6 AUG 18
NEWS
                Data available for download as a PDF in RDISCLOSURE
NEWS
     7 AUG 18
                 Simultaneous left and right truncation added to PASCAL
NEWS 8 AUG 18
                FROSTI and KOSMET enhanced with Simultaneous Left and Righ
                 Truncation
NEWS 9 AUG 18
                Simultaneous left and right truncation added to ANABSTR
NEWS 10 SEP 22 DIPPR file reloaded
NEWS 11 SEP 25
                INPADOC: Legal Status data to be reloaded
NEWS 12 SEP 29 DISSABS now available on STN
NEWS 13 OCT 10
                PCTFULL: Two new display fields added
NEWS 14 OCT 21
                BIOSIS file reloaded and enhanced
NEWS 15 OCT 28 BIOSIS file segment of TOXCENTER reloaded and enhanced
NEWS EXPRESS NOVEMBER 14 CURRENT WINDOWS VERSION IS V6.01c, CURRENT
              MACINTOSH VERSION IS V6.0b(ENG) AND V6.0Jb(JP),
              AND CURRENT DISCOVER FILE IS DATED 23 SEPTEMBER 2003
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- => s serine (w) (protease or proteinase)
 - 33 FILES SEARCHED...
 - 62 FILES SEARCHED...
 - 92 FILES SEARCHED...

```
144481 SERINE (W) (PROTEASE OR PROTEINASE)
 => s substrate (4a) (Fluorescence or Fluorescent or fret)
   24 FILES SEARCHED...
   49 FILES SEARCHED...
   74 FILES SEARCHED...
          21729 SUBSTRATE (4A) (FLUORESCENCE OR FLUORESCENT OR FRET)
 => s l1 and l2
  49 FILES SEARCHED...
            854 L1 AND L2
 => s (proteinase or protease) (4A) ((fluorescence adj resonance adj energy adj
 transfer) or fret)
   15 FILES SEARCHED...
   31 FILES SEARCHED...
   48 FILES SEARCHED...
   61 FILES SEARCHED...
   70 FILES SEARCHED...
   87 FILES SEARCHED...
             89 (PROTEINASE OR PROTEASE) (4A) ((FLUORESCENCE ADJ RESONANCE ADJ
                ENERGY ADJ TRANSFER) OR FRET)
 => s 13 and 14
 <---->User Break---->
SEARCH ENDED BY USER
=> s (PROTEINASE OR PROTEASE) (4A) ((FLUORESCENCE RESONANCE ENERGY TRANSFER) OR
FRET)
  22 FILES SEARCHED...
  43 FILES SEARCHED...
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           137 (PROTEINASE OR PROTEASE) (4A) ((FLUORESCENCE RESONANCE ENERGY
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=> s 13 and 15
  30 FILES SEARCHED...
  52 FILES SEARCHED...
  85 FILES SEARCHED...
             6 L3 AND L5
=> d 16 1-6 bib ab
     ANSWER 1 OF 6 USPATFULL on STN
L6
       2003:180876 USPATFULL
AN
ΤI
       Proteases
IN
       Yang, Junming, San Jose, CA, UNITED STATES
       Baughn, Mariah R., San Leandro, CA, UNITED STATES
       Burford, Neil, Durham, CT, UNITED STATES
       Au-Young, Janice, Brisbane, CA, UNITED STATES
       Lu, Dyung Aina M., San Jose, CA, UNITED STATES
       Reddy, Roopa, Sunnyvale, CA, UNITED STATES
       Yue, Henry, Sunnyvale, CA, UNITED STATES
       Nguyen, Daniel B., San Jose, CA, UNITED STATES
       Tang, Y. Tom, San Jose, CA, UNITED STATES
       Yao, Monique G., Mountain View, CA, UNITED STATES
       Lal, Preeti, Santa Clara, CA, UNITED STATES
PΙ
       US 2003124706 A1 20030703
ΑI
       US 2002-168425
                         A1 20020621 (10)
       WO 2000-US34811
                               20001219
DТ
       Utility
FS
       APPLICATION
```

```
94304
 CLMN
        Number of Claims: 28
 ECL
        Exemplary Claim: 1
 DRWN
        No Drawings
 LN.CNT 6542
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.
        The invention provides human proteases (PRTS) and polynucleotides which
        identify and encode PRTS. The invention also provides expression
        vectors, host cells, antibodies, agonists, and antagonists. The
        invention also provides methods for diagnosing, treating, or preventing
        disorders associated with aberrant expression of PRTS.
     ANSWER 2 OF 6 USPATFULL on STN 2003:140474 USPATFULL
AN
TI
        Device for detecting bacterial contamination and method of use
ΙN
        Sanders, Mitchell C., West Boylston, MA, UNITED STATES
                        A1
PI
       US 2003096315
                                20030522
       US 2000-201405P 20 Utility
AΙ
                                20010503 (9)
PRAI
                          20000503 (60)
DT
       Utility
FS
       APPLICATION
LREP
       HAMILTON, BROOK, SMITH & REYNOLDS, P.C., 530 VIRGINIA ROAD, P.O. BOX
       9133, CONCORD, MA, 01742-9133
CLMN
       Number of Claims: 9
ECL
       Exemplary Claim: 1
DRWN
       3 Drawing Page(s)
LN.CNT 698
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       A device and method for detecting the presence or absence of a
       prokaryotic microorganism are provided, comprising the steps of
       identifying a protein, such as a microbial-specific protease that
       characterizes the presence of a specific prokaryotic microbe and thereby
       provides a marker for that microbe; detecting the protease that is a
       marker for the presence of a specific prokaryotic microbe by cleaving a
       substrate when the protease is present; and signaling the presence of
       that protease when cleavage has occurred. More specifically, the method
       comprises identifying at least one outer membrane protein or a secreted
       protein that is unique to a particular microbial pathogen such as for
       example Listeria monocytogenes and that is substrate specific.
     ANSWER 3 OF 6 USPATFULL on STN 2003:113091 USPATFULL
L6
AN
ΤI
       Production of cultured human mast cells and basophils for high
       throughput small molecule drug discovery
       Rossi, Alexander B., San Francisco, CA, UNITED STATES
IN
PΙ
       US 2003077824 A1
                               20030424
ΑI
       US 2001-53355
                         A1
                                20011108 (10)
PRAI
       US 2001-316723P
                          20010831 (60)
DT
       Utility
FS
       APPLICATION
LREP
       Robin M. Silva, Esq., DORSEY & WHITNEY LLP, Suite3400, Four Embarcadero
       Center, San Francisco, CA, 94111-4187
CLMN
       Number of Claims: 36
       Exemplary Claim: 1
ECL
DRWN
       8 Drawing Page(s)
LN.CNT 2879
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
AΒ
       Provided are methods for producing and screening proliferated
       populations of CD34-negative progenitor cells, mucosal mast cells,
       connective tissue-type mast cells and basophil cells. The methods
       generate uniform proliferated populations of cells. The proliferated
      populations contain a uniform population of a size suitable for use in
       high throughput screening methods, for example, screening for agents
```

Incyte Genomics Inc, Legal Department, 3160 Porter Drive, Palo Alto, CA,

LREP

that alter exocytosis. The invention includes screening the proliferated populations with at least one candidate bioactive agent, and evaluating the cells to detect a cell with an altered phenotype. The invention also includes isolating a candidate bioactive agent that causes the altered phenotype. Additionally, cells formed according to the described methods are also encompassed by the invention.

```
Ь6
    ANSWER 4 OF 6 USPATFULL on STN
       2002:314644 USPATFULL
ΑN
      ASSAYS FOR APOTOSIS MODULATORS
TI
      ELLIOTT, KATHRYN J., SAN DIEGO, CA, UNITED STATES
IN
       KOUNNAS, MARIA Z., SAN DIEGO, CA, UNITED STATES
      DYER, REBECCA J., SAN DIEGO, CA, UNITED STATES
      MUNOZ, BENITO, SAN DIEGO, CA, UNITED STATES
      WAGNER, STEVEN L., SAN DIEGO, CA, UNITED STATES
                               20021128
PΙ
      US 2002177120
                         Δ1
      US 1999-326472
                         A1
                               19990604 (9)
ΔΤ
DТ
      Utility
FS
      APPLICATION
      STEPHANIE L. SEIDMAN, HELLER EHRMAN WHITE & MCAULIFFE, 4250 EXCUTIVE
LREP
      SQUARE, 7TH FLOOR, LA JOLLA, CA, 920379103
CLMN
      Number of Claims: 59
ECL
      Exemplary Claim: 1
DRWN
      2 Drawing Page(s)
LN.CNT 2250
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       Recombinant cells expressing fluorescence resonance energy transfer
       reporter polypeptides and cell-based assays for apoptosis; screening
       assays for identifying and selecting candidate compounds modulating
       apoptosis.
    ANSWER 5 OF 6 USPATFULL on STN
L6
       2002:206141 USPATFULL
AN
TI
      Fluorescent assay for proteolysis
      Benkovic, Stephen J., State College, PA, UNITED STATES
       Scott, Charles P., Narberth, PA, UNITED STATES
ΡI
      US 2002110834
                          A1.
                               20020815
AΙ
      US 2002-71468
                          A1
                               20020208 (10)
      Continuation of Ser. No. US 2000-713614, filed on 15 Nov 2000, GRANTED,
RLI
      Pat. No. US 6346924 Continuation of Ser. No. US 1997-817445, filed on 30
      Apr 1997, GRANTED, Pat. No. US 6198458
      NZ 1994-264864
PRAI
                        19941104
      NZ 1995-272778
                          19950815
      WO 1995-NZ106
                          19951016
      US 2001-267440P
                          20010208 (60)
      Utility
דת
FS
      APPLICATION
      MCKEE, VOORHEES & SEASE, P.L.C., ATTN: PENNSYLVANIA STATE UNIVERSITY,
LREP
      801 GRAND AVENUE, SUITE 3200, DES MOINES, IA, 50309-2721
      Number of Claims: 13
CLMN
ECL
      Exemplary Claim: 1
      5 Drawing Page(s)
DRWN
LN.CNT 416
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      The invention includes methods for assaying protease activity. According
ΆB
       to one aspect of the present invention provides a nucleic acid construct
      having a sequence encoding an amino terminal portion of a fluorescent
      reporter fused to a sequence encoding a substrate of a protease followed
      by a sequence encoding a carboxyl terminal portion of a fluorescent
      reporter protein. The recombinant fluorescent
       substrate is then expressed in the presence of a protease. A
      change in quenching of fluorescence in the recombinant
       substrate is then detected. The change is an indication of
      protease activity.
```

```
ANSWER 6 OF 6 WPINDEX COPYRIGHT 2003 THOMSON DERWENT on STN
    2001-265889 [27]
AN
                       WPINDEX
DNC
    C2001-080448
    New serine protease termed protease T, useful for
    treating and preventing skin flaking or imbalance of desquamation.
DC
    B04 D16
    ANDRADE-GORDON, P; DARROW, A L; QI, J; ANDRADE-GRODON, P; DARROW, A
TN
     (ORTH) ORTHO-MCNEIL PHARM INC; (ANDR-I) ANDRADE-GRODON P; (DARR-I) DARROW
PΑ
    A; (QIJJ-I) QI J
CYC 95
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    NOVELTY - A protein (I) that functions as protease T protein, is new.
          DETAILED DESCRIPTION - INDEPENDENT CLAIMS are also included for the
          (1) an isolated and purified nucleic acid molecule (II) encoding
     protease T, or its functional derivatives;
          (2) an expression vector (III), containing a nucleic acid molecule
     encoding (I);
          (3) a recombinant host cell (IV) containing (III);
          (4) a monospecific antibody (Ab) immunologically reactive with
     protease T protein;
          (5) expressing protease T in a recombinant host;
          (6) identifying compounds (C) that modulate protease T protein
     activity, by combining a modulator of protease T protein activity,
     protease T protein and a labeled substrate, and measuring a change in the
     labeled substrate;
          (7) a kit comprising a nucleic acid sequence of 1110 or 1130
     nucleotides fully defined in the specification, or their fragments;
          (8) a kit comprising a serine protease T protein
     having a sequence of 290 or 315 amino acids fully defined in the
     specification, or their fragments or derivatives; and
          (9) a pharmaceutical composition (PC) or a non-pharmaceutical
     composition (NPC), comprising (IV).
          ACTIVITY - Dermatological.
          MECHANISM OF ACTION - T serine protease
     agonist/antagonist; (claimed). No supporting data given.
          USE - (C) is useful for treating a condition mediated by protease T.
     PC is useful for treating an imbalance of desquamation, by topical
     application of PC. PC is useful as a topical skin care composition. NPC is
     useful as a laundry detergent, shampoo, hard surface cleaning
```

compositions, and dish care cleaning composition (claimed). Protease T protein is useful for treating and preventing skin flaking. NPC is also useful as skin care and hair care compositions.

useful as skin care and hair care compositions.

ADVANTAGE - Protease T is less immunogenic to sensitive individuals and it provides efficient proteolytic activity in a non-natural environment.

Dwg.0/6

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1449 et 8/19/03

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